

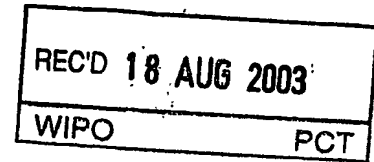


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Attestation

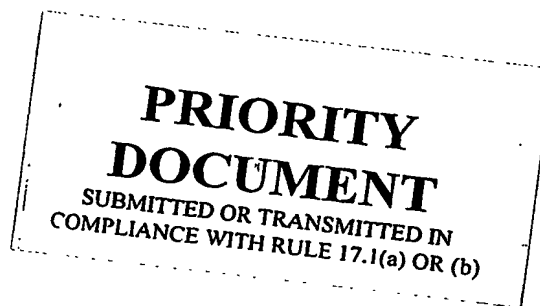
Die angehefteten Unterlagen stimmen mit der ursprünglich eingereichten Fassung der auf dem nächsten Blatt bezeichneten europäischen Patentanmeldung überein.

The attached documents are exact copies of the European patent application described on the following page, as originally filed.

Les documents fixés à cette attestation sont conformes à la version initialement déposée de la demande de brevet européen spécifiée à la page suivante.

Patentanmeldung Nr. Patent application No. Demande de brevet n°

02405633.5



Der Präsident des Europäischen Patentamts;
Im Auftrag

For the President of the European Patent Office

Le Président de l'Office européen des brevets
p.o.

R C van Dijk

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Anmeldung Nr:
Application no.: 02405633.5
Demande no:

Anmeldetag:
Date of filing: 22.07.02
Date de dépôt:

Anmelder/Applicant(s)/Demandeur(s):

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SUISSE

Bezeichnung der Erfindung/Title of the invention/Titre de l'invention:
(Falls die Bezeichnung der Erfindung nicht angegeben ist, siehe Beschreibung.
If no title is shown please refer to the description.
Si aucun titre n'est indiqué se référer à la description.)

Copolymers and their use in personal care products

In Anspruch genommene Priorität(en) / Priority(ies) claimed / Priorité(s)
revendiquée(s)
Staat/Tag/Aktenzeichen/State/Date/File no./Pays/Date/Numéro de dépôt:

Internationale Patentklassifikation/International Patent Classification/
Classification internationale des brevets:

C08F/

Am Anmeldetag benannte Vertragsstaaten/Contracting states designated at date of
filing/Etats contractants désignées lors du dépôt:

AT BE BG CH CY CZ DE DK EE ES FI FR GB GR IE IT LI LU MC NL PT SE SK TR

Copolymers and their use in personal care products

This invention relates to copolymers, their production and their use as rheology modifiers and as conditioner for water-and/or oil-based compositions, especially for water-and/or oil-based personal care products.

Rheology modifiers are used generally to adjust or modify the rheological properties of personal care compositions. Such properties include, without limitation, viscosity, flow rate, stability to viscosity change over time, and the ability to suspend particles in such personal care compositions.

Thickeners are used extensively in personal care compositions such as cosmetic and pharmaceutical formulations, to affect the aesthetics, product application and suspension and delivery of the active raw materials.

It is standard practice to include viscosifying (co-)polymers in personal care product composition in order to achieve optimum rheology characteristics. Various polymer types have been proposed for the purpose of increasing the viscosity of personal care compositions.

In US 4806345 a copolymer consisting of a quaternary ammonium salt of a cross-linked dimethylaminoethylmethacrylate and acrylamide for the use in personal care compositions is disclosed.

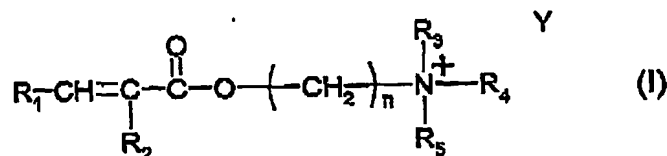
Although some copolymers used in personal care compositions described in the prior art do achieve viscosification of the composition, there is still a need to provide further improvement in rheology profile.

In addition to the rheology modifier properties mentioned in the prior art, the cationic nature of these copolymers impart substantivity to the negatively charged sites induced on the hair cuticle and stratum comeum by frictional forces, effects of UV radiation and use of chemical substances such as cleansing agents on the skin and hair. The conditioning properties of such copolymers is well known, and an increase in conditioning properties can be assessed by evaluation of the sensorial aspects of, for example, the hair, and in particular the effect on the wet and dry combing and wet and dry feel of the hair.

The aim of the present invention was to find a copolymer, which has advantages over the prior art in above-mentioned characteristics and other sensorial characteristics.

An embodiment of the present invention is a copolymer derived from the polymerization of

(a) a cationic monomer of formula (I),



wherein

R_1 is hydrogen or methyl,

R_2 is hydrogen or $\text{C}_1\text{-C}_4$ alkyl,

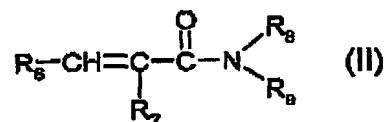
R_3 , R_4 and R_5 are independently from each other hydrogen or $\text{C}_1\text{-C}_4$ alkyl,

n is a integer from 1 – 5, and

Y is a counterion,

and

(b) a monomer of formula (II)



wherein

R_6 signifies hydrogen or methyl, and

R_7 , R_8 and R_9 signify independently from each other hydrogen or $\text{C}_1\text{-C}_4$ alkyl,

with the proviso that at least one of the substituents R_6 , R_7 , R_8 and R_9 is $\text{C}_1\text{-C}_4$ alkyl

and

(c) optionally at least one cross-linking agent, which contains at least two ethylenically unsaturated moieties.

Preferably, the copolymer consists of 20 – 95 weight-percent (wt-%) of the monomer of formula (I), more preferably of 40 – 90 wt-%, and of 5 – 50 wt-%, more preferably of 10 – 40 wt-% of the monomer of formula (II). The wt-% are based on the total weight of the polymer. The sum of the wt-% of the two components is always 100. Preferably the copolymer comprises 50 – 500 ppm, more preferably 100 – 300 ppm, of at least one cross-linking agent based on the total amount of the copolymer.

Preferably, the monomer of formula (I) is characterized in that

R_1 is hydrogen or methyl, more preferably hydrogen,

R_2 is hydrogen or methyl, more preferably hydrogen,

R_3 , R_4 and R_5 are independently from each other hydrogen or methyl, more preferably methyl,

n is an integer from 1 – 4, and

Y is Cl; Br; I; hydrogensulfate or methosulfate.

Preferably, the monomer of formula (II) is characterized in that

R_6 signifies hydrogen or methyl, more preferably hydrogen,

R_7 signifies hydrogen or methyl, more preferably hydrogen,

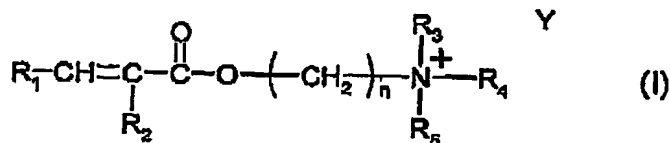
R_8 signifies hydrogen or methyl, more preferably methyl, and

R_9 signifies hydrogen or methyl, more preferably methyl,

with the proviso that at least one of the substituents R_6 , R_7 , R_8 and R_9 is methyl.

A preferred embodiment of the present invention is a copolymer derived from the polymerization of

(a) a cationic monomer of formula (I),



wherein

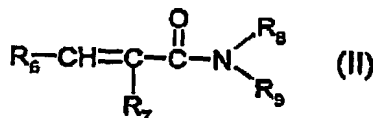
R_1 , R_2 , R_3 , R_4 and R_5 are independently from each other hydrogen or methyl,

n is 1, 2 or 3, and

Y is a counterion, preferably Cl; Br; I; hydrogensulfate or methosulfate,

and

(b) a monomer of formula (II)



wherein

R_6 signifies hydrogen or methyl, more preferably hydrogen,

R_7 signifies hydrogen or methyl, more preferably hydrogen, and

R_8 and R_9 signify independently from each other hydrogen or methyl, more preferably methyl,

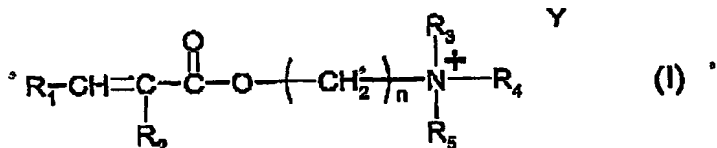
with the proviso that at least one of the substituents R_6 , R_7 , R_8 and R_9 is methyl,

and

- (c) optionally at least one cross-linking agent selected from the group of tetra allyl ammonium chloride; allyl-acrylamides and allyl-methacrylamides; bisacrylamidoacetic acid and/or N,N'-methylene-bisacrylamide, preferably tetra allyl ammonium chloride and/or N,N'-methylene-bisacrylamide.

A more preferred embodiment of the present invention is a copolymer derived from the polymerization of

- (a) 20 – 95 wt-% of a cationic monomer of formula (I), more preferably of 40 – 90 wt-% of a cationic monomer of formula (I),

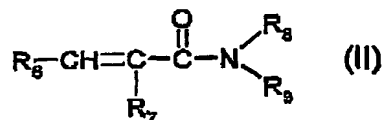


wherein

R₁, R₂, R₃, R₄ and R₅ are independently from each other hydrogen or methyl, n is 1, 2 or 3, and

Y is a counterion, preferably Cl; Br; I; hydrogensulfate or methosulfate, and

- (b) of 5 – 50 wt-%, more preferably of 10 – 40 wt-% of a monomer of formula (II)



wherein

R₆ signifies hydrogen or methyl, more preferably hydrogen,

R₇ signifies hydrogen or methyl, more preferably hydrogen, and

R₈ signifies hydrogen or methyl, more preferably methyl, and

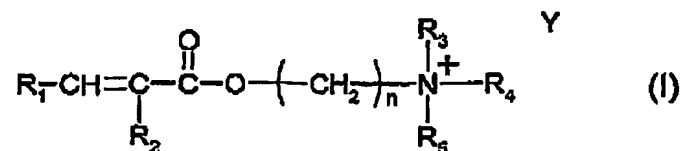
R₉ signifies hydrogen or methyl, more preferably methyl,

with the proviso that at least one of the substituents R₆, R₇, R₈ and R₉ is methyl,

- (c) of 50 – 500 ppm (based on the total amount of monomers), more preferably of 100 – 300 ppm (based on the total amount of monomers) of at least one compound of the group of tetra allyl ammonium chloride; allyl-acrylamides and allyl-methacrylamides; bisacrylamidoacetic acid and/or N,N'-methylene-bisacrylamide, more preferably tetra allyl ammonium chloride and/or N,N'-methylene-bisacrylamide.

A further embodiment is the use of a copolymer or a mixture thereof derived from the polymerization of

- (a) a cationic monomer of formula (I),



wherein

R₁ is hydrogen or methyl,

R₂ is hydrogen or C₁-C₄alkyl,

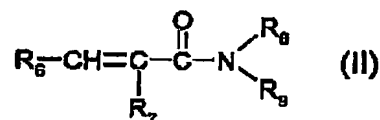
R₃, R₄ and R₅ are independently from each other hydrogen or C₁-C₄alkyl,

n is an integer from 1 – 5, and

Y is a counterion

and

- (b) a monomer of formula (II)



wherein

R₆ signifies hydrogen or methyl, and

R₇, R₈ and R₉ signify independently from each other hydrogen or C₁-C₄alkyl,

with the proviso that at least one of the substituents R₆, R₇, R₈ and R₉ is

C₁-C₄alkyl,

and

- (c) optionally at least one cross-linking agent, which contains at least two ethylenically unsaturated moieties

for water- and/or oil-based compositions, preferably for water- and/or oil-based personal care products.

A further preferred embodiment of the present invention is the use of copolymers consisting of 20 – 95 wt-% of the monomer of formula (I), more preferably of 40 – 90 wt-%, and of 5 – 50 wt-%, more preferably of 10 – 40 wt-% of the monomer of formula (II) for water- and/or oil-based compositions, preferably for water- and/or oil-based personal care products.

The wt-% are based on the total weight of the polymer.

The sum of the wt-% of the two components is always 100.

Preferably the copolymer comprises 50 – 500 ppm, more preferably 100 – 300 ppm, of at least one cross-linking agent based on the total amount of the copolymer.

A more preferred embodiment of the present invention is the use of above-mentioned copolymer wherein the monomer of formula (I) is characterized in that

R_1 is hydrogen or methyl, more preferably hydrogen,

R_2 is hydrogen or methyl, more preferably hydrogen,

R_3 , R_4 and R_5 are independently from each other hydrogen or methyl, more preferably methyl,

n is an integer from 1 – 4, and

Y is Cl; Br; I; hydrogensulfate or methosulfate,

for water- and/or oil-based compositions, preferably for water- and/or oil-based personal care compositions.

A further more preferred embodiment of the present invention is the use of above-mentioned copolymer wherein monomer of formula (II) is characterized in that

R_6 signifies hydrogen or methyl, more preferably hydrogen,

R_7 signifies hydrogen or methyl, more preferably hydrogen, and

R_8 signifies hydrogen methyl, and

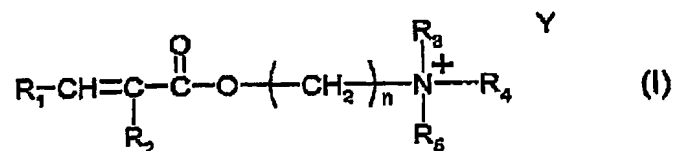
R_9 signifies hydrogen or methyl, more preferably methyl,

with the proviso that at least one of the substituents R_6 , R_7 , R_8 and R_9 is methyl,

for water- and/or oil-based compositions, preferably for water- and/or oil-based personal care compositions.

An especially preferred use is characterized in that a copolymer derived from the polymerization of

(a) a cationic monomer of formula (I),



wherein

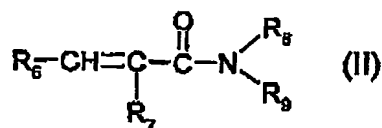
R_1 , R_2 , R_3 , R_4 and R_5 are independently from each other hydrogen or methyl,

n is 1, 2 or 3, and

Y is a counterion, preferably Cl; Br; I; hydrogensulfate or methosulfate,

and

(c) a monomer of formula (II)



wherein

R₆ signifies hydrogen or methyl, more preferably hydrogen,

R₇ signifies hydrogen or methyl, more preferably hydrogen, and

R₈ signifies hydrogen or methyl, and

R₉ signifies methyl or hydrogen, more preferably methyl.

with the proviso that at least one of the substituent R₆, R₇, R₈ and R₉ is C₁-C₄alkyl,

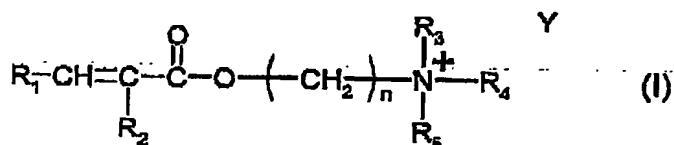
and

- (c) optionally at least one cross-linking agent selected from the group of tetra allyl ammonium chloride; allyl-acrylamides and allyl-methacrylamides; bisacrylamidoacetic acid and/or N,N'-methylene-bisacrylamide, preferably tetra allyl ammonium chloride and/or N,N'-methylene-bisacrylamide

is used for water- and/or oil-based compositions, preferably for water- and/or oil-based personal care compositions.

An very especially preferred use is characterized in that a copolymer derived from the polymerization of

- (a) 20 – 95 wt-% of a cationic monomer of formula (I), more preferably of 40 – 90 wt-% of a cationic monomer of formula (I),

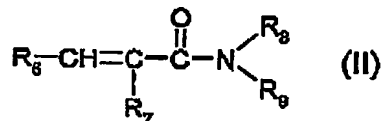


wherein

R₁, R₂, R₃, R₄ and R₅ are independently from each other hydrogen or methyl, n is 1, 2 or 3, and

Y is a counterion, preferably Cl; Br; I; hydrogensulfate or methosulfate, and

- (b) of 5 – 50 wt-%, more preferably of 10 – 40 wt-% of a monomer of formula (II)



wherein

R₆ signifies hydrogen or methyl, more preferably hydrogen,

R₇ signifies hydrogen or methyl, more preferably hydrogen, and

R₈ signifies hydrogen or methyl, and

R₉ signifies hydrogen or methyl, more preferably methyl

with the proviso that at least one of the substituent R₆, R₇, R₈ and R₉ is methyl,

and

- (c) of 50 – 500 ppm (based on the total amount of monomers), more preferably of 100 – 300 ppm (based on the total amount of monomers) of at least one compound of the group of tetra allyl ammonium chloride; allyl-acrylamides and allyl-methacrylamides; bisacrylamidoacetic acid and/or N,N'-methylene-bisacrylamide, more preferably tetra allyl ammonium chloride and/or N,N'-methylene-bisacrylamide.

The thickener-rheology modifier system of this invention is useful in personal care compositions and more particularly in hair care and skin care compositions. These compositions will generally comprise at least one cosmetically-functional agent used in an amount effective to impart desired cosmetic properties to the personal care composition. The term "cosmetically-functional agent", as used herein, means any material, compound or composition applied to the hair or skin for cosmetic application thereof. Exemplary agents include emollients, humectants, lubricants, UV-light inhibitors, preservatives, pigments, dyes, colorants, alpha-hydroxy acids, aesthetic enhancers such as starch, perfumes and fragrances, film formers (water proofing agent), antiseptics, antifungal, antimicrobial and other medicaments, solvents, surfactants, natural or synthetic polymers, hair conditioning agents and hair fixatives. Such cosmetically-functional agents include mineral oils, glycerin, beeswax, lanolin, acetylated lanolin, stearic acid, palmitic acid, cetyl alcohol, sodium salts of olefin sulfonates, various proteins, polymeric sugars, conditioning agents such as polyquaternium and hair fixatives such as poly(vinyl pyrrolidone) and N-vinyl formamide or polyvinyl formamide.

The cosmetically-functional agent may be present in the personal care composition in an amount of up to 60 wt-% (the wt-% based on the total weight of the personal care composition).

Personal care compositions are for example shampoos, bath- and shower additives, hair-care products, wax/fat compositions, liquid soaps, lotions, gels, crèmes, deodorants, stick preparations, powders, ointments, other aqueous or alcoholic or

aqueous/alcoholic solutions, for example cleaning solutions for the skin, moist cleaning sheets and oils.

Personal care compositions include a very wide range of products. Suitable products are, for example, especially the following:

- skin-care products, for example skin washing and cleansing products in the form of bars of soap or liquid soaps, syndets or washing pastes,
- bath products, for example liquid (foam baths, milks, shower products) or solid bath products, such as bath pearls and bath salts;
- skin-care products, such as skin emulsions, multiple emulsions or skin oils;
- decorative body-care products, for example face make-ups in the form of day or powder creams, face powders (loose and compressed), rouge or cream make-ups, eye-care products, for example eye shadow products, mascara, eyeliners, eye creams or eye-fix creams; lip-care products, for example lipstick, lip gloss, lip liner, nail-care products, such as nail varnish, nail varnish remover, nail hardeners or cuticle removers;
- feminine hygiene products, such as feminine hygiene washing lotions or sprays;
- foot-care products, for example foot baths, foot powders, foot creams or foot balms, special deodorants and antiperspirants or products for scrubbing off callouses;
- sunscreens, such as sun milks, lotions, creams, oils, sunblockers or tropicals, pre-sun products or after-sun products;
- suntanning products, for example self-tanning creams;
- depigmenting products, for example products for bleaching or lightening skin;
- insect repellents, for example insect oils, lotions, sprays or sticks;
- deodorants, for example deodorant sprays, non-aerosol sprays, deodorant gels, sticks or roll-ons;
- antiperspirants, for example antiperspirant sticks, creams or roll-ons;
- products for cleansing and treating impure skin, for example syndets (solid or liquid), peeling or scrubbing products or peeling masks;
- chemical depilatory products, for example depilatory powders, liquid depilatory products, creamy or pasty depilatory products, depilatory gels or aerosol foams;
- shaving products, for example shaving soap, foaming shaving creams, non-foaming shaving creams, shaving foams and gels, preshaving products for dry shaving, aftershaves or aftershave lotions;

- scents, for example perfumes (Eau de Cologne, Eau de Toilette, Eau de Parfum, Parfum de Toilette, perfume), perfume oils or perfume creams;
- products for oral and dental hygiene as well as for dentures, for example toothpastes, tooth gels, tooth powders, mouth-wash concentrates, anti-plaque mouth-washes, denture cleaning products or denture adhesion products;
- cosmetic formulations for hair treatment, for example hair washes in the form of shampoos, hair conditioners, hair-care products, for example pretreatment products, hair tonics, hair styling creams and gels, pomades, hair rinses, deep conditioning treatments, intensive hair care treatments, hair setting products, for example waving agents for perms (hot wave, mild wave, cold wave), hair straightening products, liquid hair fixatives, hair foams, hair sprays, bleaching agents, for example hydrogen peroxide solutions, bleaching shampoos, bleaching creams, bleaching powders, bleaching pastes or oils, temporary, semitemporary or permanent hair dyes, products containing self-oxidizing dyes, or natural hair dyes, such as henna or camomile.

The personal care compositions listed above can be in a very wide range of forms of presentation, for example

- in the form of liquid formulations as an oil/water (O/W) emulsion,
- in the form of a gel,
- in the form of an oil, cream, milk or lotion,
- in the form of a powder, lacquer, pellets or make-up,
- in the form of a stick,
- in the form of a spray (spray with propellant or pumping spray) or an aerosol,
- in the form of a foam, or
- in the form of a paste.

The oral hygiene composition may comprise an additional antibacterial enhancing agent, for example an anionic polymeric polycarboxylate, a dehydrated polyphosphate salt, a compound which provides a source of fluoride ions, a polishing material, including siliceous material or sodium bicarbonate, an orally acceptable vehicle, including a water-phase with humectants, thickeners, surface-active agents and a flavoring or sweetening material.

The oil phase (oil-component) can be chosen from the following substance groups without limiting the kind of lipophilic ingredient to those substances:

Fatty alcohols: Guerbet alcohols based on fatty alcohols having from 6 to 18, preferably from 8 to 10 carbon atoms including cetyl alcohol, stearyl alcohol, cetearyl alcohol, oleyl alcohol, octyldodecanol, benzoate of C_{12} - C_{15} alcohols, acetylated lanolin alcohol, etc.

Esters of fatty acids: esters of linear C_6 - C_{24} fatty acids with linear C_3 - C_{24} alcohols, esters of branched C_6 - C_{13} carboxylic acids with linear C_6 - C_{24} fatty alcohols, esters of linear C_6 - C_{24} fatty acids with branched alcohols, especially 2-ethylhexanol, esters of hydroxycarboxylic acids with linear or branched C_6 - C_{22} fatty alcohols, especially dioctyl malates, esters of linear and/or branched fatty acids with polyhydric alcohols (for example propylene glycol, dimer diol or trimer triol) and/or Guerbet alcohols, for example caproic acid, caprylic acid, 2-ethylhexanoic acid, capric acid, lauric acid, isotridecanoic acid, myristic acid, palmitic acid, palmitoleic acid, stearic acid, isostearic acid, oleic acid, elaidic acid, petroselinic acid, linoleic acid, linolenic acid, elaeostearic acid, arachidic acid, gadoleic acid, behenic acid and erucic acid and technical-grade mixtures thereof (obtained, for example, in the pressure removal of natural fats and oils, in the reduction of aldehydes from Roelen's oxosynthesis or in the dimerisation of unsaturated fatty acids) with alcohols, for example, isopropyl alcohol, caproic alcohol, capryl alcohol, 2-ethylhexyl alcohol, capric alcohol, lauryl alcohol, isotridecyl alcohol, myristyl alcohol, cetyl alcohol, palmoleyl alcohol, stearyl alcohol, isostearyl alcohol, oleyl alcohol, elaidyl alcohol, petroseliny alcohol, linoyl alcohol, linolenyl alcohol, elaeostearyl alcohol, arachidyl alcohol, gadoleyl alcohol, behenyl alcohol, erucyl alcohol and brassidyl alcohol and technical-grade mixtures thereof (obtained, for example, in the high-pressure hydrogenation of technical-grade methyl esters based on fats and oils or aldehydes from Roelen's oxo-synthesis and as monomer fractions in the dimerisation of unsaturated fatty alcohols).

Examples of such ester oils are isopropylmyristate, isopropylpalmitate, isopropylstearate, isopropyl isostearate, isopropyloleate, n-butylstearate, n-hexyllaurate, n-decylolate, isooctylstearate, iso-nonylstearate, isononyl isononanoate, 2-ethylhexylpalmitate, 2-hexyllaurate, 2-hexyldecylstearate, 2-octyldodecylpalmitate, oleylolate, oleylerucate, erucylolate, erucylrucate, cetearyl octanoate, cetyl palmitate, cetyl stearate, cetyl oleate, cetyl behenate, cetyl acetate, myristyl myristate, myristyl behenate, myristyl oleate, myristyl stearate, myristyl palmitate, myristyl lactate, propylene glycol dicaprylate/caprinate, stearyl heptanoate, diisostearyl malate, octyl hydroxystearate, etc.

Further oil components that can be used are dicarboxylic acid esters, such as diethylhexyl 2,6-naphthalate, di-n-butyl adipate, di(2-ethylhexyl)-adipate, di(2-

ethylhexyl)-succinate and diisotridecyl acetate, and also diol esters, such as ethylene glycol dioleate, ethylene glycol diisotridecanoate, propylene glycol di(2-ethylhexanoate), propylene glycol diisostearate, propylene glycol dipelargonate, butanediol diisostearate and neopentyl glycol dicaprylate. Esters of C_6 - C_{24} fatty alcohols and/or Guerbet alcohols with aromatic carboxylic acids, saturated and/or unsaturated, especially benzoic acid, esters of C_2 - C_{12} dicarboxylic acids with linear or branched alcohols having from 1 to 22 carbon atoms or polyols having from 2 to 10 carbon atoms and from 2 to 6 hydroxy groups.

Natural or synthetic triglycerides including glyceryl esters and derivatives. Di- or triglycerides, based on C_6 - C_{18} fatty acids, modified by reaction with other alcohols (caprylic/capric triglyceride, wheat germ glycerides, etc.). Fatty acid esters of polyglycerin (polyglyceryl-n such as polyglyceryl-4 caprate, polyglyceryl-2 isostearate, etc. or castor oil (*Ricinus Communis*), hydrogenated vegetable oil, sweet almond oil, wheat germ oil, sesame oil, hydrogenated cottonseed oil, coconut oil, avocado oil, corn oil, hydrogenated castor oil, shea butter, cocoa butter, soybean oil, mink oil, sunflower oil, safflower oil, macadamia nut oil, olive oil, hydrogenated tallow, apricot kernel oil, hazelnut oil, borago oil, etc.

Waxes including esters of long-chain acids and alcohols as well as compounds having wax-like properties, e.g., carnauba wax (*Copernicia Cerifera*), beeswax (white or yellow), lanolin wax, candellila wax (*Euphorbia Cerifera*), ozokerite, japan wax, paraffin wax, microcrystalline wax, ceresin, cetearyl esters wax, synthetic beeswax, etc. Also, hydrophilic waxes as Cetearyl Alcohol or partial glycerides.

Pearlescent waxes: 1,4-ethylene glycol esters, especially ethylene glycol distearate; fatty acid-alkanolamides; especially coco fatty acid diethanolamide; partial glycerides, especially stearic acid monoglyceride; esters of polyvalent, unsubstituted or hydroxy-substituted carboxylic acids with fatty alcohols having from 6 to 22 carbon atoms, especially long-chained esters of tartaric acid; fatty substances, for example fatty alcohols, fatty ketones, fatty aldehydes, fatty ethers and fatty carbonates, which in total have at least 24 carbon atoms, especially laurone and distearyl ether; fatty acids, such as stearic acid, hydroxystearic acid or behenic acid, ring-opening products of olefin epoxides having from 12 to 22 carbon atoms with fatty alcohols having from 12 to 22 carbon atoms and/or polyols having from 2 to 15 carbon atoms and from 2 to 10 hydroxy groups, and mixtures thereof.

Hydrocarbon oils: mineral oil (light or heavy), petrolatum (yellow or white), microcrystalline wax, paraffinic and isoparaffinic compounds, hydrogenated isoparaffinic molecules as polydecenes, and polybutene, hydrogenated

polyisobutene, squalane, isohexadecane, isododecane and others from plant and animal kingdom.

Silicones or siloxanes (organosubstituted polysiloxanes): dimethylpolysiloxanes, methyl/phenylpolysiloxanes, cyclic silicones, and also amino-, fatty acid-, alcohol-, polyether-, epoxy-, fluorine-, glycoside- and/or alkyl-modified silicone compounds, which at room temperature may be in either liquid or resinous form. Linear polysiloxanes: dimethicone such as Dow Corning® 200 fluid, Mirasil® DM (Rhodia), dimethiconol. Cyclic silicone fluids: cyclopentasiloxanes volatiles such as Dow Corning® 345 fluid, Silbione® grade, Abil® grade. Phenyltrimethicone ; Dow coming® 556 fluid. Also suitable are simethicones, which are mixtures of dimethicones having an average chain length of from 200 to 300 dimethylsiloxane units with hydrogenated silicates. A detailed survey by Todd *et al.* of suitable volatile silicones may in addition be found in Cosm. Toil. 91, 27 (1976).

Fluorinated or perfluorinated oils: perfluorhexane, dimethylcyclohexane, ethylcyclopentane (Flutec® grades), polyperfluoromethylisopropyl ether (Fomblin® grades)

The oil components can be used in an amount of up to 99 wt-%.

In an O/W-formulation the oil component is preferably present in an amount of from 5 wt-% to 50 wt-% and more preferably from 10 wt-% to 35 wt-%, based on the total weight of the personal care composition.

Emulsifiers. Any conventionally usable emulsifier can be used for the personal care compositions. Emulsifier systems may comprise for example:

Carboxylic acids and their salts: alcalin soap of sodium, potassium and ammonium, metallic soap of calcium or magnesium, organic basis soap such as Lauric, palmitic, stearic and oleic acid etc...

Alkyl phosphates or phosphoric acid esters: acid phosphate, diethanolamine phosphate, potassium cetyl phosphate.

Ethoxylated Carboxylic Acids or Polyethyleneglycol esters (PEG-n Acylates). Linear fatty alcohols having from 8 to 22 carbon atoms, branched from 2 to 30 mol of ethylene oxide and/or from 0 to 5 mol propylene oxide with fatty acids having from 12 to 22 carbon atoms and with alkylphenols having from 8 to 15 carbon atoms in the alkyl group. Fatty alcohol Polyglycolether such as Laureth-n, Ceteareth-n, Steareth-n,

Oleth-n. Fatty acid polyglycolether such as PEG-n Stearate, PEG-n Oleate, PEG-n Cocoate

Monoglycerides and Polyol esters. C_{12} - C_{22} fatty acid mono- and di-esters of addition products of from 1 to 30 mol of ethylene oxide with polyols

Fatty acid and polyglycerol ester such as Monostearate glycerol, diisostearoyl polyglyceryl-3-diisostearates, polyglyceryl-3-diisostearates, triglyceryl diisostearates, polyglyceryl-2-sesquiisostearates or polyglyceryl dimerates. Mixtures of compounds from a plurality of those substance classes are also suitable. Fatty acid polyglycolesters such as Monostearate diethylene glycol, Fatty acid and Polyethylene glycol esters, Fatty acid and saccharose esters such as Sucro esters, glycerol and saccharose esters such as Sucro glycerides

Sorbitol and sorbitan: Sorbitan mono- and di-esters of saturated and unsaturated fatty acids having from 6 to 22 carbon atoms and ethylene oxide addition products

Polysorbate-n series, Sorbitan esters such as Sesquiisostearate, sorbitan, PEG-(6)-isostearate sorbitan, PEG-(10)-Sorbitan laurate, PEG-17- dioleate sorbitan.

Glucose derivatives: C_8 - C_{22} alkyl-mono and oligo-glycosides and ethoxylated analogues with glucose being preferred as the sugar component. O/W emulsifiers such as Methyl Gluceth-20 sesquistearate, Sorbitan Stearate/Sucrose Cocoate, Methyl Glucose Sesquistearate, Cetearyl alcohol/Cetearyl glucoside. W/O emulsifiers such as Methyl glucose Dioleate/ Methyl glucose isostearate.

Sulfates and sulfonated derivatives: Dialkylsulfosuccinates (DOSS: Dioctyl succinate), alkyl lauryl sulfonate, linear sulfonated paraffins, sulfonated tetrapropylene sulfonate, sodium lauryl sulfates, ammonium and ethanolamine lauryl sulfates, lauryl ether sulfates, sodium laureth sulfates, sulfosuccinates, acetyl isothionates, alkanolamide sulfates such as Taurines, Methyl taurines, Imidazole sulfates.

Amine derivatives: amine salts, ethoxylated amines such as Oxide amine, with chains containing an heterocycle such as alkyl imidazolines, pyridine derivatives, isoquinoteines, cetyl pyridinium chloride, cetyl pyridinium bromure, quaternary ammonium such as cetyltrimethylbromure ammonium bromure (CTBA), Stearylalkonium.

Amide derivatives. alkanolamides such as acylamide DEA, ethoxylated amides, such as PEG-n acylamide, oxydeamide.

Polysiloxane/Polyalkyl/Polyether Copolymers and derivatives: dimethicone, copolyols, silicone polyethylene Oxide copolymer, silicone glycol copolymer.

Propoxylated or POE-n ethers (Meroxapols), Polaxamers or poly(oxyethylene)m-block-poly(oxypropylene)n-block(oxyethylene)

Zwitterionic surfactants that carry at least one quaternary ammonium group and at least one carboxylate and/or sulfonate group in the molecule. Zwitterionic surfactants that are especially suitable are the so-called betaines, such as N-alkyl-N,N-dimethylammonium glycinate, for example cocoalkyldimethylammonium glycinate, N-acylaminoethyl-N,N-dimethylammonium glycinate, for example cocoacylaminoethyl-N,N-dimethylammonium glycinate, and 2-alkyl-3-carboxymethyl-3-hydroxyethylimidazolines each having from 8 to 18 carbon atoms in the alkyl or acyl group and also cocoacylaminoethylhydroxyethyl-carboxy-methylglycinate, N-alkylbetaine, N-alkylaminobetaines.

Alkylimidazolines, alkylpeptides, lipoaminoacides

Self emulsifying bases (. K:F:DePolo – A short textbook of cosmetology, Chapter 8, Table 8-7, p250-251):

Non ionic bases such as PEG-6 Beeswax (and) PEG-6 Stearate (and) polyglyceryl – 2 isostearate [Apifac], Glyceryl stearate (and) PEG-100 stearate. [Arlacel 165], PEG-5 Glyceryl stearate [Arlatone 983 S], Sorbitan oleate (and) Polyglyceryl-3 Ricinoleate.[Arlacel 1689], Sorbitan Stearate and sucrose cocoate [Arlatone 2121], Glyceryl stearate and laureth-23 [Cerasynth 945], Cetearyl alcohol and ceteth-20 [Cetomacrogol Wax], Cetearyl alcohol and Polysorbate 60 and PEG-150 and stearate-20[Polawax GP 200, Polawax NF], Cetearyl alcohol and cetearyl polyglucoside [Emulgade PL 1618], Cetearyl alcohol and cetareth-20 [Emulgade 1000NI, Cosmowax], Cetearyl alcohol and PEG-40 castor oil [Emulgade F Special], Cetearyl Alcohol and PEG-40 Castor Oil and Sodium Cetearyl Sulfate [Emulgade F], Stearyl Alcohol and Steareth-7 and Steareth-10 [Emulgator E 2155], Cetearyl Alcohol and Steareth-7 and steareth-10 [Emulsifying wax U.S.N.F], Glyceryl-stearate and PEG-75 stearate [Gelot 64], Propylene Glycol ceteth-3 Acetate [Hetester PCS], Propylene Glycol isoceth-3 Acetate [Hetester PHA], Cetearyl alcohol and Ceteth-12 and oleth-12 [Lanbriol Wax N 21], PEG –6 Stearate and PEG-32 Stearate [Tefose 1500], PEG-6 Stearate and ceteth-20 and steareth-20 [Tefose 2000], PEG-6 Stearate and ceteth-20 and Glyceryl Stearate and steareth-20 [Tefose 2561], Glyceryl Stearate and cetareth-20 [Teginacid H, C, X].

Anionic alkaline bases such as PEG-2 Stearate SE, Glyceryl stearate SE [Monelgine, Cutina KD], Propylene glycol stearate [Tegin P].

Anionic acid bases such as Cetearyl Alcohol and Sodium cetearyl sulfate [Lanette N, Cutina LE, Crodacol GP], Cetearyl Alcohol and sodium Lauryl Sulfate [Lanette W], Trilaneth-4 Phosphate and glycol stearate and PEG-2 Stearate [Sedefos 75], Glyceryl stearate and sodium lauryl Sulfate [Teginacid Special].

Cationic acid bases such as cetearyl Alcohol and cetrimonium bromide.

The emulsifiers may be used in an amount of, for example, from 1 wt-% to 30 wt-%, especially from 4 wt-% to 20 wt-% and preferably from 5 wt-% to 10 wt-%, based on the total weight of the composition.

When formulated in O/W emulsions, the preferably amount of such emulsifier system could represent 5% to 20% of the oil phase.

Adjuvants and additives. The personal care compositions, for example creams, gels, lotions, alcoholic and aqueous/alcoholic solutions, emulsions, wax/fat compositions, stick preparations, powders or ointments, may in addition contain, as further adjuvants and additives, mild surfactants, super-fatting agents, consistency regulators, additional thickeners, polymers, stabilizers, biogenic active ingredients, deodorizing active ingredients, anti-dandruff agents, film formers, swelling agents, further UV light-protective factors, antioxidants, hydrotropic agents, preservatives, insect repellents, self-tanning agents, solubilizers, perfume oils, colourants, bacteria-inhibiting agents and the like.

The adjuvants and additives may optionally be present in the personal care composition in an amount of, for example, from 0.1 wt-% to 25 wt-% based on the total weight of the composition.

Super-fatting agents. Substances suitable for use as super-fatting agents are, for example, lanolin and lecithin and also polyethoxylated or acrylated lanolin and lecithin derivatives, polyol fatty acid esters, monoglycerides and fatty acid alkanolamides, the latter simultaneously acting as foam stabilizers.

Surfactants. Examples of suitable mild surfactants, that is to say surfactants especially well tolerated by the skin, include fatty alcohol polyglycol ether sulfates, monoglyceride sulfates, mono- and/or di-alkyl sulfosuccinates, fatty acid isothionates, fatty acid sarcosinates, fatty acid taurides, fatty acid glutamates, α -olefin sulfonates, ethercarboxylic acids, alkyl oligoglucosides, fatty acid glucamides, alkylamidobetaines and/or protein fatty acid condensation products, the latter preferably being based on wheat proteins.

Consistency regulators / Additional Thickeners and Rheology modifiers. As additional thickeners and rheology modifiers, there come into consideration the groups of silicium dioxide, magnesium silicates, aluminium silicates, polysaccharides or derivatives thereof for example hyaluronic acid, xanthan gum, guar-guar, agar-

agar, alginates, Carraghenan, gellan, pectines, or modified cellulose such as hydroxycellulose, hydroxypropylmethylcellulose. In addition polyacrylates or homopolymer of reticulated acrylic acids and polyacrylamides, e.g. the Carbopol range (e.g. Carbopol types 980, 981, 1382, ETD 2001, ETD2020, Ultrez 10; INCI: Carbomer) or Salcare range such as Salcare SC80 (Stearth-10 Allyl Ether/Acrylates Copolymer), Salcare SC81 (Acrylates copolymer), Salcare SC91 and Salcare AST (Sodium Acrylates Copolymer/PPG-1 trideceth-6), Sepigel 305 (Polyacrylamide/laureth-7), Simulgel NS and Simulgel EG (Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer), Stablen 30 (Acrylates/Vinyl Isodecanoate Crosspolymer), Pemulen TR-1 (Acrylates/C₁₀-C₃₀ Alkyl Acrylate Crosspolymer), Luvigel EM (Sodium Acrylates Copolymer), Aculyn 28 (Acrylates/Beheneth-25 Methacrylate Copolymer), etc.

Polymers. Suitable cationic polymers are, for example, cationic cellulose derivatives, for example a quaternized hydroxymethyl cellulose obtainable under the name Polymer JR 400[®] from Amerchol, cationic starches, copolymers of diallylammonium salts and acrylamides, quaternized vinylpyrrolidone/vinyl imidazole polymers, for example Luviquat[®] (BASF), condensation products of polyglycols and amines, quaternized collagen polypeptides, for example lauryldimonium hydroxypropyl hydrolyzed collagen (Lamequat[®]/L/Grünau), quaternised wheat polypeptides, polyethyleneimine, cationic silicone polymers, for example amidomethicones, copolymers of adipic acid and dimethylaminohydroxypropyldiethylenetriamine (Cartaretin[®]/Clariant), copolymers of acrylic acid with dimethyldiallylammonium chloride (Merquat[®] 550/Chemviron), polyaminopolyamides, as described, for example, in FR-A-2 252 840, and the cross-linked water-soluble polymers thereof, cationic chitin derivatives, for example of quaternised chitosan, optionally distributed as microcrystals; condensation products of dihaloalkyls, for example dibromobutane, with bisdialkylamines, for example bisdimethylamino-1,3-propane, cationic guar gum, for example Jaguar[®] C-17, Jaguar[®] C-16 from Celanese, quaternised ammonium salt polymers, for example Mirapol[®] A-15, Mirapol[®] AD-1, Mirapol[®] AZ-1 from Miranol. As anionic, zwitterionic, amphoteric and non-ionic polymers there come into consideration, for example, vinyl acetate/crotonic acid copolymers, vinylpyrrolidone/vinyl acrylate copolymers, vinyl acetate/butyl maleate/isobornyl acrylate copolymers, methyl vinyl ether/maleic anhydride copolymers and esters thereof, uncrosslinked polyacrylic acids and polyacrylic acids cross linked with polyols, acrylamidopropyltrimethylammonium chloride/acrylate copolymers, octyl

acrylamide/methyl methacrylate/tert-butylaminoethyl methacrylate/2-hydroxypropyl methacrylate copolymers, polyvinylpyrrolidone, vinylpyrrolidone/vinyl acetate copolymers, vinylpyrrolidone/dimethyl-aminoethyl methacrylate/vinyl caprolactam terpolymers and also optionally derivatised cellulose ethers and silicones. Furthermore the polymers as described in EP 1093796 (pages 3 - 8, paragraphs 17 - 68) may be used.

Biogenic active ingredients. Biogenic active ingredients are to be understood as meaning, for example, tocopherol, tocopherol acetate, tocopherol palmitate, ascorbic acid, deoxyribonucleic acid, retinol, bisabolol, allantoin, phytantriol, panthenol, AHA acids, amino acids, ceramides, pseudoceramides, essential oils, plant extracts and vitamin complexes.

Deodorizing active ingredients. As deodorizing active ingredients there come into consideration, for example, antiperspirants, for example aluminium chlorohydrates (see J. Soc. Cosm. Chem. 24, 281 (1973)). Under the trade mark Locron® of Clariant, there is available commercially, for example, an aluminium chlorohydrate corresponding to formula $Al_2(OH)_5Cl \times 2.5 H_2O$, the use of which is especially preferred (see J. Pharm. Pharmacol. 26, 531 (1975)). Besides the chlorohydrates, it is also possible to use aluminium hydroxyacetates and acidic aluminium/zirconium salts. Esterase inhibitors may be added as further deodorizing active ingredients. Such inhibitors are preferably trialkyl citrates, such as trimethyl citrate, tripropyl citrate, triisopropyl citrate, tributyl citrate and especially triethyl citrate (Hydagen® CAT, Henkel KGaA, Düsseldorf/GER), which inhibit enzyme activity and hence reduce odour formation. Further substances that come into consideration as esterase inhibitors are sterol sulfates or phosphates, for example lanosterol, cholesterol, campesterol, stigmasterol and sitosterol sulfate or phosphate, dicarboxylic acids and esters thereof, for example glutaric acid, glutaric acid monoethyl ester, glutaric acid diethyl ester, adipic acid, adipic acid monoethyl ester, adipic acid diethyl ester, malonic acid and malonic acid diethyl ester and hydroxycarboxylic acids and esters thereof, for example citric acid, malic acid, tartaric acid or tartaric acid diethyl ester. Antibacterial active ingredients that influence the germ flora and kill or inhibit the growth of sweat-decomposing bacteria can likewise be present in the preparations (especially in stick preparations). Examples include chitosan, phenoxyethanol and chlorhexidine gluconate. 5-Chloro-2-(2,4-dichlorophenoxy)-phenol (Irgasan®, Ciba Specialty Chemicals Inc.) has also proved especially effective.

Anti-dandruff agents. As anti-dandruff agents there may be used, for example, climbazole, octopirox and zinc pyrithione. Customary film formers include, for example, chitosan, microcrystalline chitosan, quaternised chitosan, polyvinylpyrrolidone, vinylpyrrolidone/vinyl acetate copolymers, polymers of quaternary cellulose derivatives containing a high proportion of acrylic acid, collagen, hyaluronic acid and salts thereof and similar compounds.

Antioxidants.

The personal care product can optionally contain one or more antioxidants. Any common antioxidant can be used.

Typical examples of such antioxidants are 4,4'-di- α -cumyl-diphenylamine, mono- and dialkylated tert-butyl/tert-octyl-diphenylamines, n-octadecyl 3,5-di-tert-butyl-4-hydroxyhydrocinnamate, Tetradibutyl pentaerythrityl-4-hydroxyhydrocinnamate, neopentane-tetrayl tetrakis(3,5-di-tert-butyl-4-hydroxyhydrocinnamate), di-n-octadecyl 3,5-di-tert-butyl-4-hydroxybenzylphosphonate, 1,3,5-tris(3,5-di-tert-butyl-4-hydroxybenzyl)isocyanurate, thioldiethylene bis(3,5-di-tert-butyl-4-hydroxyhydrocinnamate), 1,3,5-trimethyl-2,4,6-tris(3,5-di-tert-butyl-4-hydroxybenzyl)benzene, 3,6-dioxaoctamethylene bis(3-methyl-5-tert-butyl-4-hydroxyhydrocinnamate), 2,6-di-tert-butyl-p-cresol, 2,2'-ethylidene-bis(4,6-di-tert-butylphenol), 1,3,5-tris(2,6-dimethyl-4-tert-butyl-3-hydroxybenzyl) isocyanurate, 1,1,3-tris(2-methyl-4-hydroxy-5-tert-butylphenyl)butane, 1,3,5-tris[2-(3,5-di-tert-butyl-4-hydroxyhydrocinnamoyloxy)ethyl] isocyanurate, 3,5-di-(3,5-di-tert-butyl-4-hydroxybenzyl)mesitol, hexamethylene bis(3,5-di-tert-butyl-4-hydroxyhydrocinnamate), 1-(3,5-di-tert-butyl-4-hydroxyanilino)-3,5-di(octylthio)-s-triazine, N,N'-hexamethylene-bis(3,5-di-tert-butyl-4-hydroxyhydrocinnamamide), calcium bis(ethyl 3,5-di-tert-butyl-4-hydroxybenzylphosphonate), ethylene bis[3,3-di(3-tert-butyl-4-hydroxyphenyl)butyrate], octyl 3,5-di-tert-butyl-4-hydroxybenzylmercaptoacetate, bis(3,5-di-tert-butyl-4-hydroxyhydrocinnamoyl)-hydrazide, N,N'-bis[2-(3,5-di-tert-butyl-4-hydroxyhydrocinnamoyloxy)-ethyl]oxamide, and N,N-dialkylhydroxylamine prepared from di(hydrogenated tallow)amine by direct oxidation.

Further suitable antioxidants are amino acids (e.g. glycine, histidine, tyrosine, tryptophan) and derivatives thereof, imidazoles (e.g. urocanic acid) and derivatives thereof, peptides, such as D,L-carnosine, D-carnosine, L-carnosine and derivatives thereof (e.g. anserine), carotinoids, carotenes (e.g. β -carotene, β -carotene, lycopene) and derivatives thereof, chlorogenic acid and derivatives thereof, lipoic acid and derivatives thereof (e.g. dihydrolipoic acid), aurothioglyucose, propylthiouracil and

other thiols (e.g. thioredoxin, glutathione, cysteine, cystine, cystamine and the glycosyl, N-acetyl, methyl, ethyl, propyl, amyl, butyl, lauryl, palmitoyl, oleyl, linoleyl, cholesteryl and glyceryl esters thereof) and also salts thereof, dialauryl thiodipropionate, distearyl thiodipropionate, thiodipropionic acid and derivatives thereof (esters, ethers, peptides, lipids, nucleotides, nucleosides and salts) and also sulfoximine compounds (e.g. buthionine sulfoximines, homocysteine sulfoximine, buthionine sulfones, penta-, hexa-, hepta-thionine sulfoximine), also (metal) chelating agents (e.g. α -hydroxy fatty acids, palmitic acid phytic acid, lactoferrin), α -hydroxy acids (e.g. citric acid, lactic acid, malic acid), humic acid, bile acid, bile extracts, bilirubin, biliverdin, EDTA, EGTA and derivatives thereof, unsaturated fatty acids and derivatives thereof (e.g. linolenic acid, linoleic acid, oleic acid), folic acid and derivatives thereof, ubiquinone and ubiquinol and derivatives thereof, vitamin C and derivatives (e.g. ascorbyl palmitate, magnesium ascorbyl phosphate, ascorbyl acetate), tocopherols and derivatives (e.g. vitamin E acetate), vitamin A and derivatives (e.g. vitamin A palmitate) and also coniferyl benzoate of benzoin resin, rutinic acid and derivatives thereof, α -glycosylrutin, ferulic acid, furfurylidene glucitol, camosine, butyl hydroxytoluene, butyl hydroxyanisole, nordihydroguaiaretic acid, trihydroxybutyrophenone, uric acid and derivatives thereof, mannose and derivatives thereof, superoxide dismutase, N-[3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionyl]-sulfanilic acid (and salts thereof, for example the disodium salts), zinc and derivatives thereof (e.g. ZnO, ZnSO₄), selenium and derivatives thereof (e.g. selenium methionine), stilbene and derivatives thereof (e.g. stilbene oxide, trans-stilbene oxide) and the derivatives (salts, esters, ethers, sugars, nucleotides, nucleosides, peptides and lipids) of HALS ("Hindered Amine Light Stabilizers") compounds may also be mentioned.

The amount of antioxidants present is usually from 0.001 wt-% to 30 wt-%, preferably from 0.01 wt-% to 3 wt-%, based on the weight of the personal care product.

Hydrotropic agents. To improve the flow behaviour it is also possible to employ hydrotropic agents, for example ethoxylated or non ethoxylated mono-alcohols, diols or polyols with a low number of C-atoms or their ethers (e.g. ethanol, isopropanol, 1,2-dipropanediol, propyleneglycol, glycerine, ethylene glycol, ethylene glycol monoethylether, ethylene glycol monobutylether, propylene glycol monomethylether, propylene glycol monoethylether, propylene glycol monobutylether, diethylene glycol monomethylether, diethylene glycol monoethylether, diethylene glycol monobutylether and similar products). The polyols that come into consideration for

that purpose have preferably from 2 to 15 carbon atoms and at least two hydroxy groups. The polyols may also contain further functional groups, especially amino groups, and/or may be modified with nitrogen. Typical examples are as follows: glycerol, alkylene glycols, for example ethylene glycol, diethylene glycol, propylene glycol, butylene glycol, hexylene glycol and also polyethylene glycols having an average molecular weight of from 100 to 1000 Dalton; technical oligoglycerol mixtures having an intrinsic degree of condensation of from 1.5 to 10, for example technical diglycerol mixtures having a diglycerol content of from 40 wt-% to 50 wt-%; methylol compounds, such as, especially, trimethylolmethane, trimethylolpropane, trimethylolbutane, pentaerythritol and dipentaerythritol; lower alkyl-glucosides, especially those having from 1 to 8 carbon atoms in the alkyl radical, for example methyl and butyl glucoside; sugar alcohols having from 5 to 12 carbon atoms, for example sorbitol or mannitol; sugars having from 5 to 12 carbon atoms, for example glucose or saccharose; amino sugars, for example glucamine; dialcohol amines, such as diethanolamine or 2-amino-1,3-propanediol.

Preservatives and Bacteria-inhibiting agents. Suitable preservatives include, for example, methyl-, ethyl-, propyl-, butyl- parabens, Benzalkonium chloride, 2-Bromo-2-nitro-propane-1,3-diol, Dehydroacetic acid, Diazolidinyl Urea, 2-Dichloro-benzyl alcohol, DMDM hydantoin, Formaldehyde solution, Methylidibromoglutanitrile, Phenoxyethanol, Sodium Hydroxymethylglycinate, Imidazolidinyl Urea and Triclosan and further substance classes listed in the following reference: K.:F.DePolo – A short textbook of cosmetology, Chapter 7, Table 7-2, 7-3, 7-4 and 7-5, p210-219.

Bacteria-inhibiting agents. Typical examples of bacteria-inhibiting agents are preservatives that have a specific action against gram-positive bacteria, such as 2,4,4'-trichloro-2'-hydroxydiphenyl ether, chlorhexidine (1,6-di(4-chlorophenyl-biguanido)hexane) or TCC (3,4,4'-trichlorocarbanilide). A large number of aromatic substances and ethereal oils also have antimicrobial properties. Typical examples are the active ingredients eugenol, menthol and thymol in clove oil, mint oil and thyme oil. A natural deodorizing agent of interest is the terpene alcohol farnesol (3,7,11-trimethyl-2,6,10-dodecatrien-1-ol), which is present in lime blossom oil. Glycerol monolaurate has also proved to be a bacteriostatic agent. The amount of the additional bacteria-inhibiting agents present is usually from 0.1 wt-% to 2 wt-%, based on the solids content of the preparations.

Perfume oils. There may be mentioned as perfume oils mixtures of natural and/or synthetic aromatic substances. Natural aromatic substances are, for example, extracts from blossom (lilies, lavender, roses, jasmine, neroli, ylang-ylang), from stems and leaves (geranium, patchouli, petitgrain), from fruit (aniseed, coriander, caraway, juniper), from fruit peel (bergamot, lemons, oranges), from roots (mace, angelica, celery, cardamom, costus, iris, calmus), from wood (pinewood, sandalwood, guaiacum wood, cedarwood, rosewood), from herbs and grasses (tarragon, lemon grass, sage, thyme), from needles and twigs (spruce, pine, Scots pine, mountain pine), from resins and balsams (galbanum, elemi, benzoin, myrrh, olibanum, opoponax). Animal raw materials also come into consideration, for example civet and castoreum. Typical synthetic aromatic substances are, for example, products of the ester, ether, aldehyde, ketone, alcohol or hydrocarbon type. Aromatic substance compounds of the ester type are, for example, benzyl acetate, phenoxyethyl isobutyrate, p-tert-butylcyclohexyl acetate, linalyl acetate, dimethylbenzylcarbonyl acetate, phenylethyl acetate, linalyl benzoate, benzyl formate, ethylmethylphenyl glycinate, allylcyclohexyl propionate, styralyl propionate and benzyl salicylate. The ethers include, for example, benzyl ethyl ether; the aldehydes include, for example, the linear alkanals having from 8 to 18 hydrocarbon atoms, citral, citronellal, citronellyl oxyacetaldehyde, cyclamen aldehyde, hydroxycitronellal, lillial and bourgeonal; the ketones include, for example, the ionones, α,α -isomethylionone and methyl cedryl ketone; the alcohols include, for example, anethol, citronellol, eugenol, isoeugenol, geraniol, linalool, phenyl ethyl alcohol and terpinol; and the hydrocarbons include mainly the terpenes and balsams. It is preferable, however, to use mixtures of various aromatic substances that together produce an attractive scent. Ethereal oils of relatively low volatility, which are chiefly used as aroma components, are also suitable as perfume oils, e.g. sage oil, camomile oil, clove oil, melissa oil, oil of cinnamon leaves, lime blossom oil, juniper berry oil, vetiver oil, olibanum oil, galbanum oil, labolanum oil and lavandin oil. Preference is given to the use of bergamot oil, dihydromyrcenol, lillial, lylal, citronellol, phenyl ethyl alcohol, α,α -hexyl cinnamaldehyde, geraniol, benzyl acetone, cyclamen aldehyde, linalool, boisambrene forte, ambroxan, indole, hedione, sandelice, lemon oil, tangerine oil, orange oil, allyl amyl glycolate, cyclovertal, lavandin oil, muscatel sage oil, α -damascone, bourbon geranium oil, cyclohexyl salicylate, vertofix coeur, iso-E-Super, Fixolide NP, evernyl, lraldein gamma, phenylacetic acid, geranyl acetate, benzyl acetate, rose oxide, romillat, irotyl and floramat alone or in admixture with one another.

Colourants. There may be used as colourants the substances that are suitable and permitted for cosmetic purposes, as compiled, for example, in the publication "Kosmetische Färbemittel" of the Farbstoffkommission der Deutschen Forschungsgemeinschaft, Verlag Chemie, Weinheim, 1984, pages 81 to 106. The colourants are usually used in concentrations of from 0.001 wt-% to 0.1 wt-%, based on the total mixture.

Other adjuvants. It is furthermore possible for the personal care composition to contain, as adjuvants, anti-foams, such as silicones, *structurants*, such as maleic acid, *solubilisers*, such as ethylene glycol, propylene glycol, glycerol or diethylene glycol, opacifiers, such as latex, styrene/PVP or styrene/acrylamide copolymers, complexing agents, such as EDTA, NTA, alaninediacetic acid or phosphonic acids, *propellants*, such as propane/butane mixtures, N₂O, dimethyl ether, CO₂, N₂ or air, so-called coupler and developer components as oxidation dye precursors, *reducing agents*, such as thioglycolic acid and derivatives thereof, thiolactic acid, cysteamine, thiomalic acid or mercaptoethanesulfonic acid, or oxidizing agents, such as hydrogen peroxide, potassium bromate or sodium bromate.

There come into consideration as insect repellents, for example, N,N-diethyl-m-toluamide, 1,2-pentanediol or insect repellent 3535; suitable self-tanning agents are, for example, dihydroxyacetone and/or erythrulose or dihydroxy acetone and/or dihydroxy acetone precursors as described in WO 01/85124 and/or erythrulose.

Ultraviolet Light Absorbers

Ultraviolet Light Absorbers (UV absorbers) are employed in cosmetics to protect the product from chemical or physical deterioration induced by ultraviolet light. Sunscreen Agents are OTC drug ingredients, which protect the skin from ultraviolet light. UV absorbers, like sunscreen agents, have the ability to convert incident ultraviolet radiation into less damaging infrared radiation (heat). Suitable UV absorbers are, for example:

Acetaminosalol, Allantoin PABA, Benzalphenalide, Benzophenone, Benzophenone-1, Benzophenone-2, Benzophenone-3, Benzophenone-4, Benzophenone-5, Benzophenone-6, Benzophenone-7, Benzophenone-8, Benzophenone-9, Benzophenone-10, Benzophenone-11, Benzophenone-12, Benzotriazolyl Dodecyl p-Cresol, 3-Benzylidene Camphor, Benzylidenecamphor Hydrolyzed Collagen Sulfonamide, Benzylidene Camphor Sulfonic Acid, Benzyl Salicylate, Bis-Ethylhexyloxyphenol Methoxyphenyl Triazine, Borelone, Bumetizole, Butyl

Methoxydibenzoylmethane, Butyl PABA, Callophyllum Inophyllum Seed Oil, Camellia Sinensis Leaf Extract, Carotenoids, Ceria/Silica, Ceria/Silica Talc, Cinoxate, DEA-Methoxycinnamate, Dibenzoxazolyl Naphthalene, Di-t-Butyl Hydroxybenzylidene Camphor, Diethylhexyl Butamido Triazone, Diethylhexyl 2,6-Naphthalate, Digalloyl Trioleate, Diisopropyl Methyl Cinnamate, 1-(3,4-Dimethoxyphenyl)-4,4-Dimethyl-1,3-Pentanediene, Dimethyl PABA Ethyl Cetearyldimonium Tosylate, Dimorpholinopyridazinone, Diphenyl Carbomethoxy Acetoxy Naphthopyran, Disodium Bisethylphenyl Triaminotriazine Stilbenedisulfonate, Disodium Distyrylbiphenyl Disulfonate, Disodium Phenyl Dibenzimidazole Tetrasulfonate, Drometrizole, Drometrizole Trisiloxane, Esculin, Ethyl Dihydroxypropyl PABA, Ethyl Diisopropylcinnamate, Ethylhexyl Dimethoxybenzylidene Dioxoimidazolidine Propionate, Ethylhexyl Dimethyl PABA, Ethylhexyl Ferulate, Ethylhexyl Methoxycinnamate, Ethylhexyl Salicylate, Ethylhexyl Triazone, Ethyl Methoxycinnamate, Ethyl PABA, Ethyl Urocanate, Etocrylene, Ferulic Acid, 4-(2-Beta-Glucopyranosiloxy) Propoxy-2-Hydroxybenzophenone, Glyceryl Ethylhexanoate Dimethoxycinnamate, Glyceryl PABA, Glycol Salicylate, Hexanediol salicylate, Homosalate, Hydrolyzed Lupine Protein, Isoamyl p-Methoxycinnamate, Isopentyl Trimethoxycinnamate Trisiloxane, Isopropylbenzyl Salicylate, Isopropyl Dibenzoylmethane, Isopropyl Methoxycinnamate, Menthyl Anthranilate, Menthyl Salicylate, 4-Methylbenzylidene Camphor, Methylene Bis-Benzotriazolyl Tetramethylbutylphenol, Octocrylene, Octrizole, PABA, PEG-25 PABA, Pentyl Dimethyl PABA, Phenylbenzimidazole Sulfonic Acid, Pinus Pinaster Bark Extract, Polyacrylamidomethyl Benzylidene Camphor, Polysilicone-15, Potassium Methoxycinnamate, Potassium Phenylbenzimidazole Sulfonate, Red Petrolatum, Sodium Benzotriazolyl Butylphenol Sulfonate, Sodium Isoferulate, Sodium Phenylbenzimidazole Sulfonate, Sodium Urocanate, Spirulina Platensis Powder, TEA-Phenylbenzimidazole Sulfonate, TEA-Salicylate, Terephthalylidene Dicamphor Sulfonic Acid, Tetrabutyl Phenyl Hydroxybenzoate, Titanium Dioxide, Tocotrienols, TriPABA Panthenol, Urocanic Acid, VA/Crotonates/Methacryloxybenzophenone-1 Copolymer and Vitis Vinifera (Grape) Seed Extract.

Polymeric beads or hollow spheres as SPF enhancers. The combination of the UV-absorbers described above, with SPF enhancers, such as non-active ingredients like Styrene/acrylates copolymer, silica beads, spheroidal magnesium silicate, spherical polyamide powder such as n-lactam polymer (Orgasol® range, Elf Atochem) cross linked Polymethylmethacrylates (PMMA; Micoparl M305 Seppic), can maximize better the UV protection of the sun products. Holosphere additives

(Sunspheres® ISP, Silica Shells Kobo.) deflect radiation and the effective path length of the photon is therefore increased.(EP0893119). Some beads, as mentioned previously, provide a soft feel during spreading. Moreover, the optical activity of such beads, e.g.Micropearl M305, cans modulate skin shine by eliminating reflection phenomena and indirectly may scatter the UV light.

When formulated in O/W emulsions, the preferably amount of such SPF enhancers should represent 1% to 10% of the total amount of the personal care composition.

A typical O/W- based personal care composition comprises:

0.5 – 10 wt-% of at least one of the inventive copolymer

2 – 25 wt-% of at least one oil-component,

0 – 25wt-% of at least one adjuvant and/or additive,

water up to 100 wt-%.

A typical oil-based personal care composition comprises

0.5 – 10 wt-% of at least one of the inventive copolymer

50 – 99 wt-% of at least one oil-component,

0 – 25 wt-% of at least one adjuvant and/or additive.

An soap has, for example, the following composition:

0.1 to 5% wt-% of at least one inventive copolymer

0.3 to 1% wt-% of titanium dioxide

1 to 10% wt-% of stearic acid

to 100 wt-% of soap base, for example the sodium salts of tallow fatty and coconut fatty acid or glycerols.

A shampoo has, for example, the following composition:

0.1 to 5% wt-% of at least one inventive copolymer

12.0% wt-% of sodium laureth-2-sulfate,

4.0% wt-% of cocamidopropylbetaine,

3.0% wt-% of NaCl and

water to 100 wt-%.

A deodorant has, for example, the following composition:

0.1 to 5% of at least one inventive copolymer

60% by weight of ethanol,

0.3% by weight of perfume oil and

water to 100%.

The new cationic polymers are obtainable by conventional polymerization processes.

The following examples are set forth merely to exemplify the invention and are not intended to limit the metes and bounds of the invention, which is set forth by the claims appended hereto.

Parts and percentages are by weight. The temperatures are stated in degrees Celsius.

Synthesis of the liquid dispersion copolymer*

This example illustrates the preparation of a suitable cationic liquid dispersion copolymer.

An 'aqueous phase' of water soluble components is prepared by admixing together the following components:

0.29 parts	Citric acid-1-hydrate.
0.18 parts	of a 40% solution of penta sodium diethylene triamine penta acetic
40.77 parts	of water
0.01 parts	of methylene-bis-acrylamide
11.75 parts	of N,N-dimethyl acrylamide (99%)
47.0 parts	of Methyl Chloride quaternised dimethylaminoethylmethacrylate.

An 'oil phase' is prepared by admixing together the following components:

1.7 parts	of sorbitan tri-oleate
3.25 parts	of a polymeric stabilizer
63.25 parts	of a medicinal grade mineral oil
24.7 parts	of Isopar [®] G (dearomatised hydrocarbon solvent; Isopar is a trademark of the Exxon Mobil Corporation)

The two phases are mixed together in a ratio of 1 part oil phase to 1.25 parts aqueous phase under high shear to form a water-in-oil emulsion.

The resulting water-in-oil emulsion is transferred to a reactor equipped with nitrogen sparge tube, stirrer and thermometer. The emulsion is purged with nitrogen to remove oxygen.

Polymerization is effected by addition of a redox couple of sodium metabisulphite and tertiary butyl hydroperoxide.

After the isotherm is completed addition is made of a free radical initiator (Vazo® 67, trademark of DuPont de Nemours and Company)) and the emulsion held at 85°C for 75 minutes.

Vacuum distillation is carried out to remove water and volatile solvent to give a final product of 50% polymer solids.

To this product addition is made of 8 parts (by weight of final product) of a fatty alcohol alkoxyate (PPG1-trideceth 6).

Formulation Examples

Example 1: Facial Moisturizer

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	2.00
Coco-caprylate / Caprate	2.50
Squalane	2.00
Hexyl Laurate	2.00
Ethylhexyl Palmitate	2.00
Dimethicone	2.50
Ethylhexyl Methoxycinnamate	5.00
Parfum	0.20
Preservative	0.20

Example 2: Body Moisturizer

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	1.00
Stearyl Alcohol	5.00
Cetyl Alcohol	5.00
Dimethicone	5.00
Cetearyl Stearate	2.00
Glycerin	2.00
Propylene Glycol	2.00
Parfum	0.20
Preservative	0.20

Example 3: Spray Moisturizer

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	1.50
Cyclomethicone	3.00
Hydrogenated Polydecene	5.00
Isostearyl Lactate	1.50
Sodium Hyaluronate	1.00
Glyceryl Myristate	1.00
Parfum	0.20
Preservative	0.20

Example 4: Leave-On Conditioner

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	1.50
Propylene Glycol	2.00
Glycerin	2.00
Dimethicone Copolyol	2.00
Preservative	0.25
Parfum	0.30
Ethylhexyl Methoxycinnamate	2.00

Example 5: Silicone Conditioner

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	2.00
Cyclopentasiloxane (and) Dimethiconol	2.00
Cyclomethicone	2.00
Ceteareth-5	0.75
Preservative	0.20
Dimethicone PEG-8 Meadowfoamate	0.50
Parfum	0.20

Example 6: Rinse-Off Conditioner

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	2.00
Decyl Oleate	2.00
Helianthus Annuus	2.50
Dimethicone (and) Dimethiconol	2.50
Preservative	0.20
Parfum	0.30
CI 18965	0.02
Sodium Benzotriazolyl Butylphenol Sulfonate (and) Buteth-3 (and) Tributyl Citrate	0.20

Example 7: Sunless Tanning Cream with Sunscreen

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	2.00
Ethylhexyl Methoxycinnamate	5.00
Dihydroxyacetone	3.00
Methylene bis-Benzotriazolyl Tetramethyl Butylphenol	3.00
Paraffinum Liquidum	7.50
Preservative	0.50
Glycerin	2.00
Parfum	0.50

Example 8: Moisturizing Lipstick

Ingredient	Amount (wt-%)
Ricinus Communis	25.00
Euphorbia Cerifera	5.40
Copernicia Cerifera	4.00
Ozokerite	5.00
Hydrogenated Lanolin	11.10
Microcrystalline Wax	4.50
Polymer of the Present Invention	2.25
Octyldodecanol	6.60
Isocetyl Palmitate	5.00
Beeswax	2.00
Cetearyl Alcohol	20.00
Preservative	0.10
Tetradibutyl Pentaerythrityl Hydroxyhydrocinnamate	0.05
Pigment	9.00

Example 9: Moisturizing Soap Base

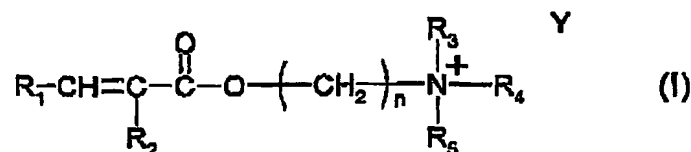
Ingredient	Amount (wt-%)
Polymer of the Present Invention	1.00
Sodium Tallowate (and) Sodium Cocoate	98.10
Tetrasodium EDTA	0.10
Titanium Dioxide	0.10
Tetradibutyl Pentaerythrityl Hydroxyhydrocinnamate	0.05
Parfum	0.50

Example 10: Anti-Acne Skin Cream

Ingredient	Amount (wt-%)
Aqua	To 100
Polymer of the Present Invention	4.00
Alcohol	5.00
Isocetyl Palmitate	2.00
Salicylic Acid	2.00
Paraffinum Liquidum	1.00
Preservative	0.50
Glycerin	2.00
Parfum	0.50

Claims

1. A copolymer derived from the polymerization of
 (a) a cationic monomer of formula (I),



wherein

R₁ is hydrogen or methyl,

R₂ is hydrogen or C₁-C₄alkyl,

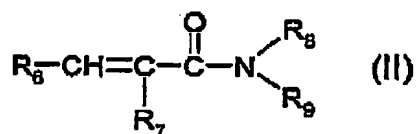
R₃, R₄ and R₅ are independently from each other hydrogen or C₁-C₄alkyl,

n is a integer from 1 - 5, and

Y is a counterion,

and

- (b) a monomer of formula (II)



wherein

R₆ signifies hydrogen or methyl, and

R₇, R₈ and R₉ signify independently from each other hydrogen or C₁-C₄alkyl,

with the proviso that at least one of the substituents R₆, R₇, R₈ and R₉ is

C₁-C₄alkyl,

and

- (c) optionally at least one cross-linking agent, which contains at least two ethylenically unsaturated moieties.

2. A copolymer according to Claim 1 characterized in that it consists of
 20 - 95 wt-% of the monomer of formula (I) and of
 5 - 50 wt-% of the monomer of formula (II).
3. A copolymer according to Claim 1 or 2 characterized in that it consists of
 40 - 90 wt-% of the monomer of formula (I) and of
 10 - 40 wt-% of the monomer of formula (II).

4. A copolymer according to anyone of the preceding claims characterized in that the copolymer comprises 50 – 500 ppm, preferably 100 – 300 ppm of at least one cross-linking agent based on the total amount of the copolymer.

5. A copolymer according to anyone of the preceding claims characterized in that

R_1 is hydrogen or methyl, more preferably hydrogen,

R_2 is hydrogen or methyl, more preferably hydrogen,

R_3 , R_4 and R_5 are independently from each other hydrogen or methyl, more preferably methyl,

n is an integer from 1 – 4, and

Y is Cl; Br; I; hydrogensulfate or methosulfate.

6. A copolymer according to anyone of the preceding claims characterized in that

R_6 signifies hydrogen or methyl, more preferably hydrogen,

R_7 signifies hydrogen or methyl, more preferably hydrogen, and

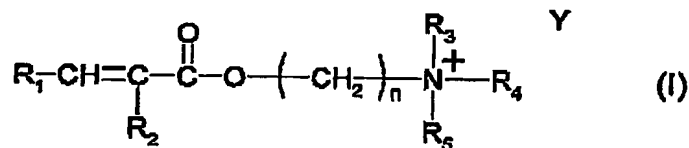
R_8 signifies hydrogen or methyl, and

R_9 signifies hydrogen or methyl, more preferably methyl,

with the proviso that at least one of the substituents R_6 , R_7 , R_8 and R_9 is methyl.

7. A copolymer according to Claim 1 derived from the polymerization of

(a) a cationic monomer of formula (I),



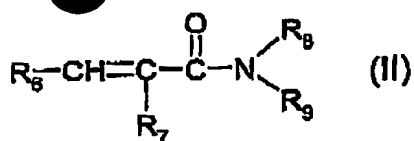
wherein

R_1 , R_2 , R_3 , R_4 and R_5 are independently from each other hydrogen or methyl,

n is 1, 2 or 3, and

Y is a counterion, preferably Cl; Br; I; hydrogensulfate or methosulfate, and

(b) a monomer of formula (II)



wherein

R_6 signifies hydrogen or methyl, more preferably hydrogen,

R_7 signifies hydrogen or methyl, more preferably hydrogen, and

R_8 signifies hydrogen or methyl, more preferably methyl, and

R_9 signifies hydrogen or methyl, more preferably methyl,

with the proviso that at least one of the substituents R_6 , R_7 , R_8 and R_9 is methyl,

and

- (c) optionally at least one cross-linking agent selected from the group of tetra allyl ammonium chloride; allyl-acrylamides and allyl-methacrylamides; bisacrylamidoacetic acid and/or N,N'-methylene-bisacrylamide, preferably tetra allyl ammonium chloride and/or N,N'-methylene-bisacrylamide.

8. A copolymer according to Claim 7 derived from the polymerization of 20 – 95 wt-% of a cationic monomer of formula (I), more preferably of 40 – 80 wt-% of a cationic monomer of formula (I),
and
of 5 – 50 wt-%, more preferably of 10 – 40 wt-% of a monomer of formula (II)
and
of 50 – 500 ppm (based on the total amount of monomers), more preferably of 100 – 300 ppm (based on the total amount of monomers) of at least one compound of the group of tetra allyl ammonium chloride; allyl-acrylamides and allyl-methacrylamides; bisacrylamidoacetic acid and/or N,N'-methylene-bisacrylamide, more preferably tetra allyl ammonium chloride and/or N,N'-methylene-bisacrylamide.
9. Use of a copolymer according to anyone of the preceding Claims for water- and/or oil-based compositions, preferably for water- and/or oil-based personal care compositions.
10. An oil/water- based personal care composition comprises:
0.5 – 10 wt-% of at least one copolymer according to Claim 1 - 8

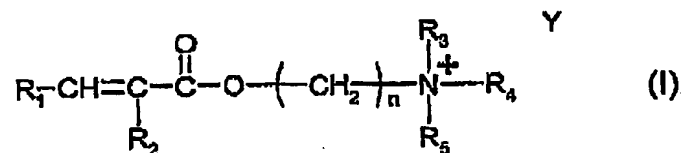
2 – 25 wt-% of at least one oil-component,
0 – 25wt-% of at least one adjuvant and/or additive,
water up to 100 wt-%.

11. A typical oil-based personal care composition comprises
0.5 – 10 wt-% of at least one copolymer according to Claim 1 - 8
50 – 99 wt-% of at least one oil-component,
0 – 25 wt-% of at least one adjuvant and/or additive.
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Abstract

The present invention relates to a copolymer derived from the polymerization of

- a) a cationic monomer of formula (I),



wherein

R₁ is hydrogen or methyl,

R₂ is hydrogen or C₁-C₄alkyl,

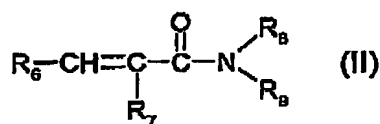
R₃, R₄ and R₅ are independently from each other hydrogen or C₁-C₄alkyl,

n is a integer from 1 - 5, and

Y is a counterion,

and

- b) a monomer of formula (II)



wherein

R₆ signifies hydrogen or methyl, and

R₇, R₈ and R₉ signify independently from each other hydrogen or C₁-C₄alkyl, with the proviso that at least one of the substituents R₆, R₇, R₈ and R₉ is C₁-C₄alkyl and

- c) optionally at least one cross-linking agent, which contains at least two ethylenically unsaturated moieties,

as well as to their use in personal care product and to the personal care products.

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